RN 128273-78-9 CAPLUS
CN Thiourea, N-[[4-[(bromodifluoromethyl)thio]phenyl]methyl]-N'-(2,2,2-trifluoroethyl)- (9CI) (CA INDEX NAME)

[--00000106]

RN 128273-79-0 CAPLUS
CN Thiourea, N-[[4-(heptafluoropropyl)phenyl]methyl]-N'-(2,2,2-trifluoroethyl)- (9CI) (CA INDEX NAME)

[--00000107]

RN 128273-80-3 CAPLUS
CN Thiourea, N-[[4-[3,3,4,4,5,5,5-heptafluoro-2,2-bis(trifluoromethyl)pentyl]phenyl]methyl]-N'-(2,2,2-trifluoroethyl)- (9CI) (CA INDEX NAME)

[--00000108]

RN 128273-81-4 CAPLUS
CN Thiourea, N-[[4-(tridecafluorohexyl)phenyl]methyl]-N'-(2,2,2-trifluoroethyl)- (9CI) (CA INDEX NAME)

[--00000109]

RN 128273-84-7 CAPLUS
CN Thiourea, N-[[4-[[3-chloro-5-(trifluoromethyl)-2-pyridinyl]oxy]phenyl]methyl]-N'-(2,2,2-trifluoroethyl)- (9CI) (CA INDEX NAME)

[--00000110]

2/9 - (C) FILE CAPLUS
STN CA Caesar accession number : 1553
AN - 1996:137693 CAPLUS
DN - 124:165248
TI - Aryl antiinflammatory compounds, their preparation, and their activity
IN - Adams, Jerry Leroy; Hall, Ralph Floyd
PA - SmithKline Beecham Corp., USA
SO - PCT Int. Appl., 46 pp.
CODEN: PIXXD2
DT - Patent
LA - English
FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PN -	WO9533458	A	19951214	WO 1995-US6961	19950602
	W: JP, US				
	RW: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE				

PR - US 1994-252718 19940602
OS - MARPAT 124:165248
AB - The invention relates to the novel compds. and pharmaceutical compns. of I [R1 = SO3H, S(O)n-C1-4 alkyl; n = 0-2; R2 = H, halo, (substituted) C1-8 alkyl, C1-8 alkoxy; m = 1, 2; R3 = C(O)R7, C(S)R7; R4, R8, R9 = H, C1-4 alkyl; R5 = H, halo, CF3, Me, (CH2)tC(O)2R8, (CH2)tOH; t = 0-2; R6 = H, halo; R7 = (substituted) aryl, (substituted) aryl-C1-2 alkyl, (substituted) C1-8 alkyl, NR9R10; R10 = (substituted) aryl, (substituted) aryl-C1-2 alkyl, (substituted) C1-8 alkyl, or R9NR10 form 5- to 7-membered (un)satd. ring with optional addnl. heteroatom of O/N or S; X = O, S; with provisions] and pharmaceutically acceptable salts thereof. The invention also relates to a method of treating or reducing inflammation in a mammal in need thereof, which comprises

of I. Prepn. of selected compds. of the invention is described. Compds. of the invention demonstrated phospholipase A2 inhibition, generally at 50 .mu.M levels.

AN - 1996:137693 CAPLUS

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PR - US 1994-252718 19940602

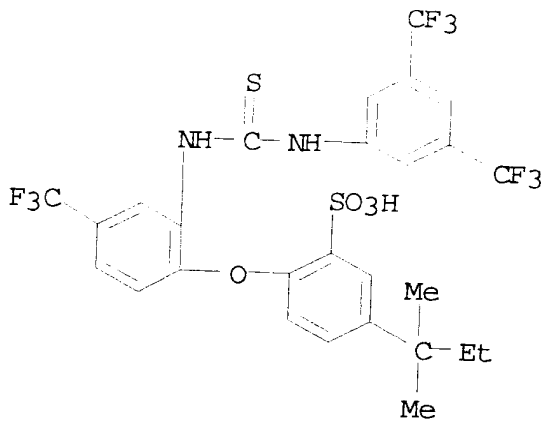
OS - MARPAT 124:165248

IT - ---174083-14-8---
RL: THU (Therapeutic use); BIOL (Biological study); USES (Uses)
(aryl antiinflammatory compd. prepn. and activity)

RN - 174083-14-8 CAPLUS

CN - Benzenesulfonic acid, 2-[2-[[[[3,5-bis(trifluoromethyl)phenyl]amino]thioxomethyl]amino]-4-(trifluoromethyl)phenoxy]-5-(1,1-dimethylpropyl)- (9CI) (CA INDEX NAME)

[--00000018]



Marker: 00000018